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## Amendments to Claims

Claim 1 (Previously Presented) A method for the expression of a coding region of interest in a Bacillus sp comprising:

- a) providing a transformed Bacillus sp cell having a chimeric gene comprising the promoter region of a Bacillus gene operably linked to a coding region of interest expressible in a Bacillus sp, wherein the promoter region is of a Bacillus subtilis yva WXY gene; and
- b) growing the transformed Bactllus sp cell of step (a) in the absence of oxygen wherein the chimeric gene of step (a) is expressed.

Claim 2 (Previously Presented) A method for the expression of a coding region of interest in a Bacillus sp comprising:

- a) providing a transformed Bacillus sp cell having a chimeric gene comprising the promoter region of a Bacillus gene operably linked to a coding region of interest expressible in a Bacillus sp, wherein the promoter region is of a Bacillus subtilis yvaWXY gene;
- b) growing the transformed Bacillus sp cell of step (a) in the presence of oxygen whereby the cell density is increased; and
- c) at a time prior to about T0, removing oxygen from the transformed Bacillus sp cell of step (b) whereby the chimeric gene is expressed.

Claim 3 (Original) A method according to Claim 2 wherein after step (c) oxygen is re-supplied to the transformed Bacillus sp cell.

Claim 4 (Previously Presented) A method according to either of Claims 1 or 2 wherein the promoter region is a promoter region for driving expression of a Bacillus gene contained in a nucleic acid fragment as set forth in SEQ ID NO:8.

## Claims 5-7 (Canceled).

Claim 8 (Previously Presented) A method for the expression of a coding region of interest in a Bacillus sp comprising:

a) providing a transformed Bacillus sp cell having a chimeric gene comprising the promoter region of a Bacillus gene operably linked to a coding region of interest expressible in a Bacillus sp, wherein the promoter region is of a Bacillus subtilis yvaWXY gene; and

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b) growing the transformed *Bacillus sp* cell of step (a) in the presence of oxygen until the cell reaches about TO of the stationary phase wherein the chimeric gene of step (a) is expressed.

Claim 9 (Previously Presented) A method according to Claim 8 wherein the promoter region is a promoter region for driving expression of a *Bacillus* gene contained in a nucleic acid fragment as set forth in SEQ ID NO:8.

## Claims 10-13 (Canceled).

Claim 14 (Previously presented) A method according to any of Claims 1, 2 or 3 wherein the expression of the chimeric gene is regulated at TO of the stationary phase.

Claim 15 (Previously Presented) A method according to any one of Claims 1, 2, 3, 4, and 8, wherein the Bacillus sp cell is selected from the species consisting of Bacillus subtilis, Bacillus thuringiensis, Bacillus anthracis, Bacillus cereus, Bacillus brevis, Bacillus megaterium, Bacillus intermedius, Bacillus thermoamyloliquefaciens, Bacillus amyloliquefaciens, Bacillus circulans, Bacillus licheniformis, Bacillus macerans, Bacillus sphaericus, Bacillus stearothermophilus, Bacillus laterosporus, Bacillus acidocaldarius, Bacillus pumilus, and Bacillus pseudofirmus.

Claim 16 (Previously presented) A method according to any one of Claims 1, 2, 3, 4, and 8, wherein the coding region of interest is selected from the group consisting of crtE crtB, pds, crtD, crtL, crtZ, crtX crtO, phaC, phaE, efe, pdc, adh, genes encoding limonene synthase, pinene synthase, bornyl synthase, phelandrene synthase, cineole synthase, sabinene synthase, and taxadiene synthase.

Claims 17-28 (Canceled).